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THYROID CARCINOMA

- Major subtypes of thyroid carcinomas are
 - Papillary carcinoma (80% to 85%)
 - Follicular carcinoma (10% to 15%)
 - Poorly differentiated and anaplastic (undifferentiated) carcinoma (<5%)
 - Medullary carcinoma (5%)

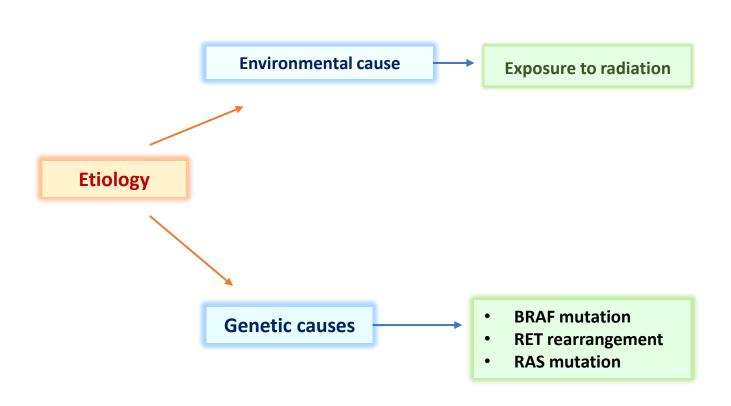


 Malignant epithelial tumor of thyroid with cells showing distinctive nuclear features and follicular cell differentiation



- Age can occur in both children and adults (median age 50 years)
- Sex predominance in females (female to male ratio 3:1)

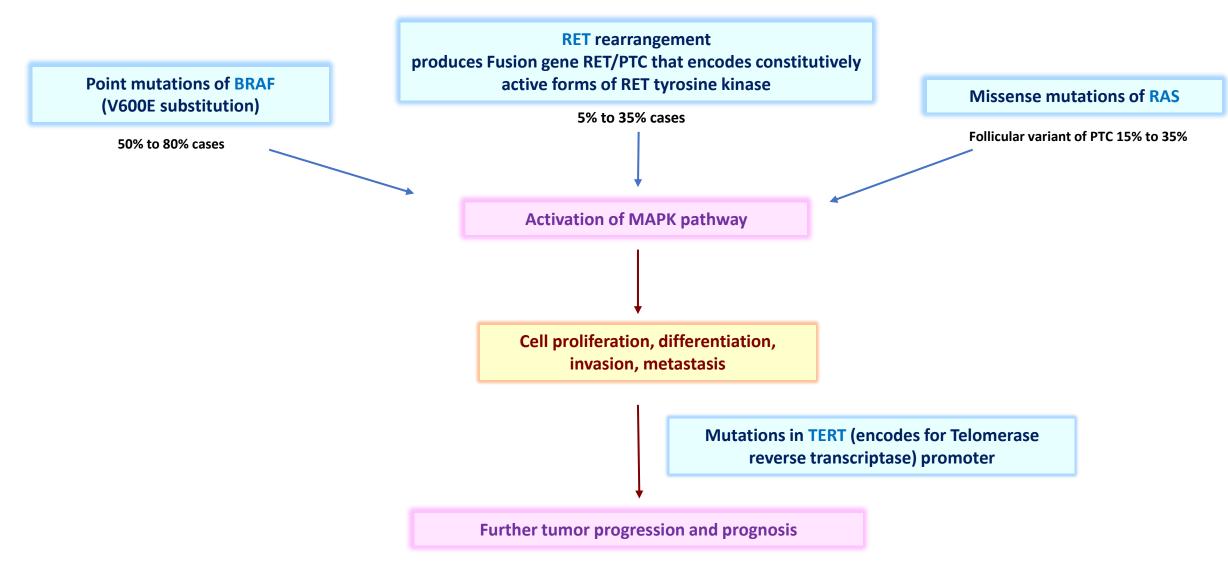




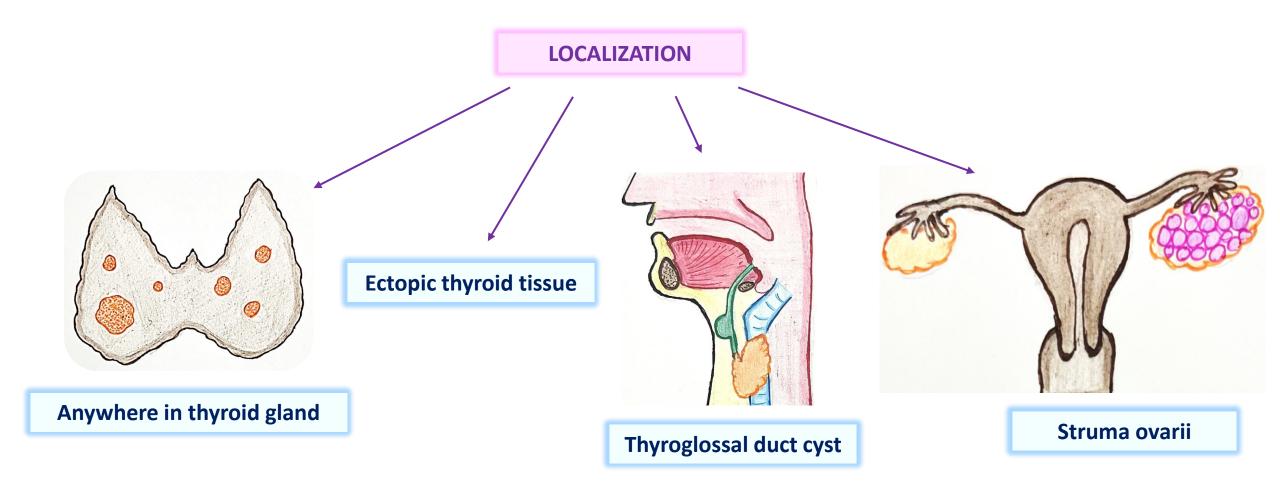
Risk factors (relationship with PTC is not clear) Obesity **Diabetes Smoking Alcohol consumption Dietary nitrates Dietary iodine excess**



Genetic alterations









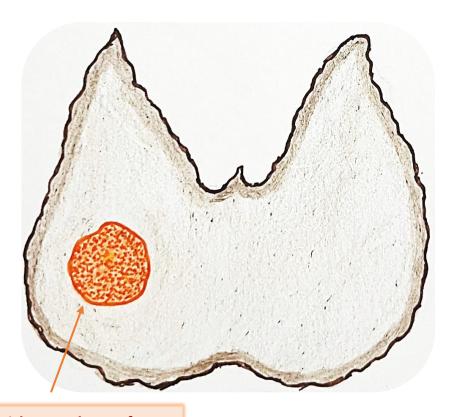
Clinical features

- Presents as asymptomatic thyroid nodule with or without involvement of cervical lymph nodes
- Hoarseness of voice and dysphagia occurs if recurrent laryngeal neve is involved
- Thyroid function test no diagnostic utility and can be normal
- Thyroid scan (123 I) usually presents as cold nodule (hypo functioning) (rarely presents as hyperfunctioning hot nodule)
- Ultrasound hypoechoic or isoechoic nodule, microcalcifications and disorganized internal vascularity



Gross:

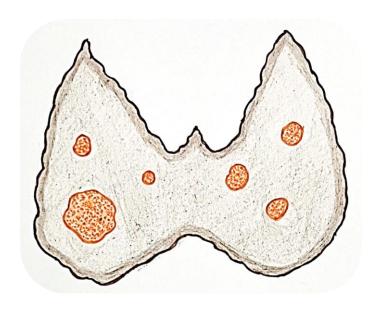
- Tumor is gray white with irregular borders and firm consistence
- Cut section granular white with specks of calcifications. Bone formation can also be seen
- Size < 1mm to several centimetres
- May have variation depending upon variant of PTC



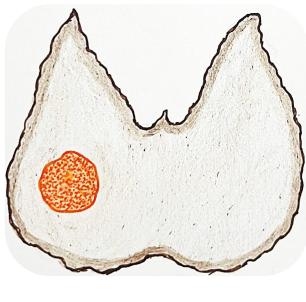
Lesion with granular surface



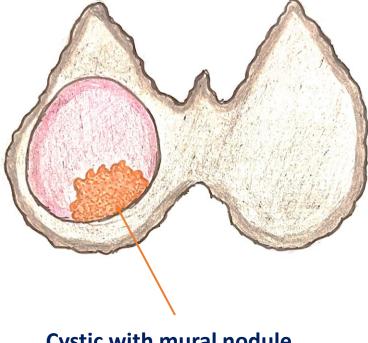
Clinical presentation



multifocal (can be due to intraglandular lymphatic spread)



unifocal



Cystic with mural nodule



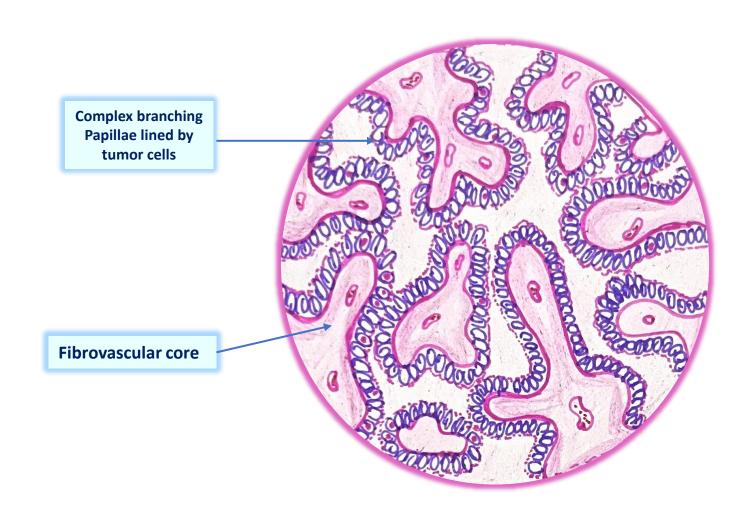
Precursor lesions recognized are

- Papillary microcarcinoma as precursor for conventional papillary thyroid carcinoma
- Non-invasive follicular thyroid neoplasm with papillary like nuclear features-
 - Low grade neoplasm with minimal risk of recurrence
 - Precursor to invasive encapsulated follicular variant of papillary thyroid carcinoma



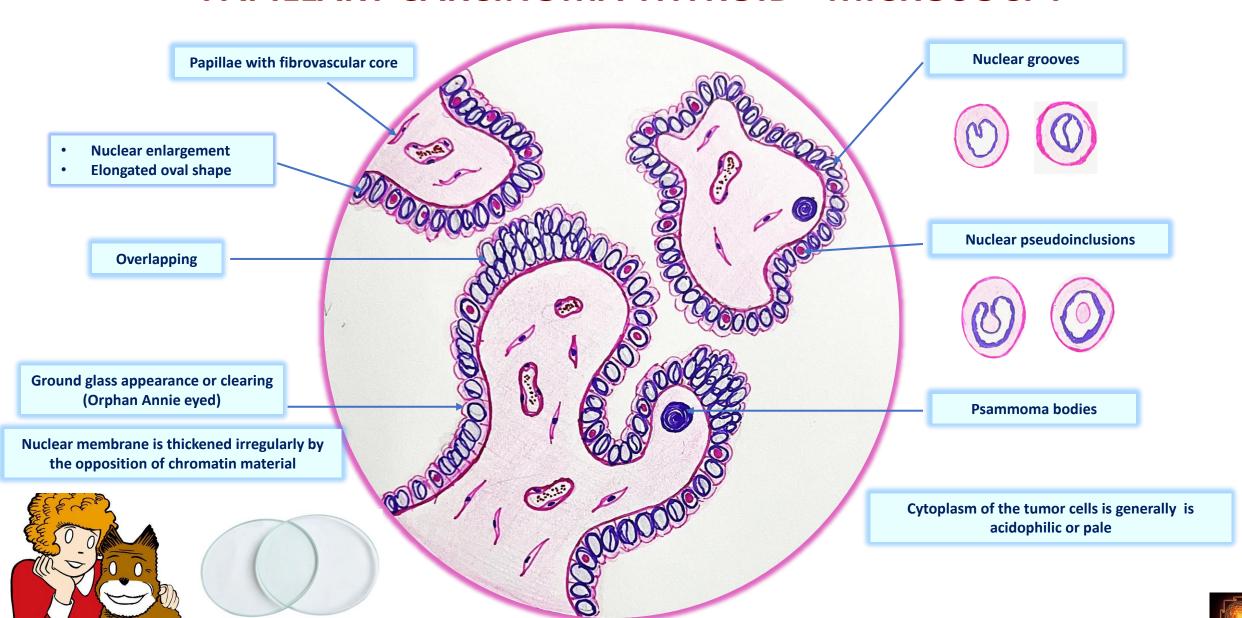
Microscopy:

- Tumor is composed of papillary structures with complex branching papillae
- Papillae have fibrovascular core and are lined by cells showing loss of polarity and with pale or eosinophilic cytoplasm
- Squamous metaplasia is also seen (20 to 40% cases)



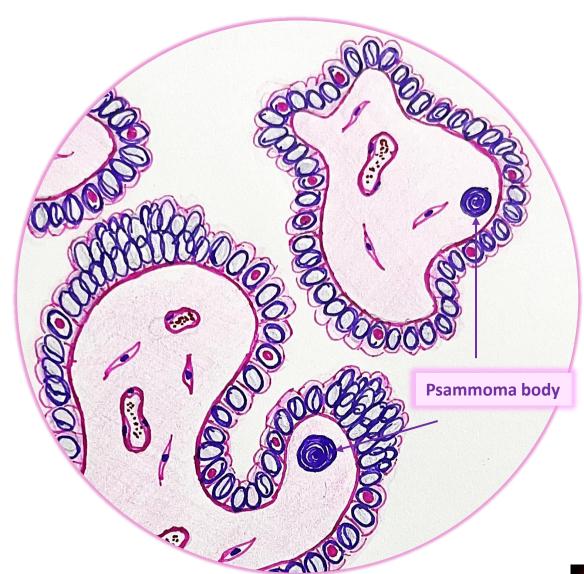


PAPILLARY CARCINOMA THYROID - MICROSOCPY



Microscopy

- Psammoma bodies may be present (50% of cases). They are rounded and laminated concentrically deposited calcium around the necrotic tumor cells (dystrophic calcification)
- They are present with in lymphatic spaces or with in tumor stroma
- Intratumoral sclerosis and peritumoral lymphocytic infiltration are found





Papillary microcarcinoma

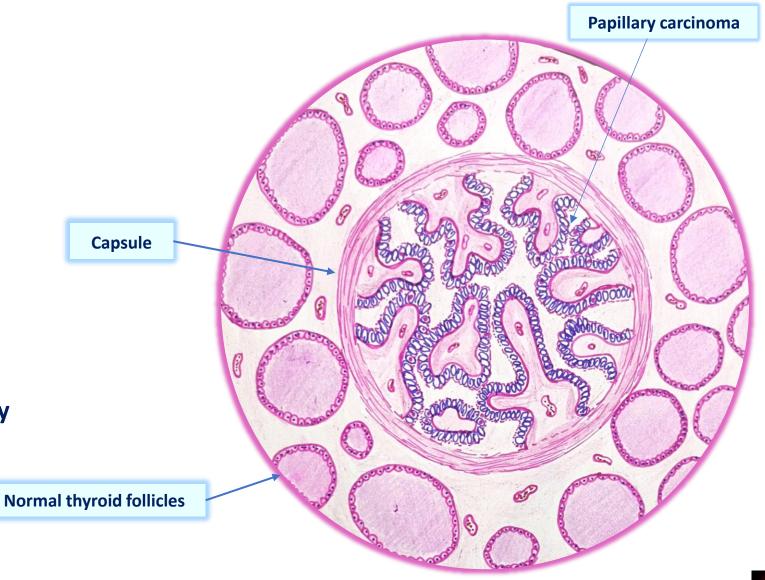
- m/s less than 1cm in diameter
- Also called occult sclerosing carcinoma, occult papillary carcinoma
- Microscopically presents as irregular scar like configuration with neoplastic elements at periphery and other elements are entrapped in the centre
- Some tumors are completely capsulated and may be focally calcified
- Prognosis excellent





Encapsulated variant-

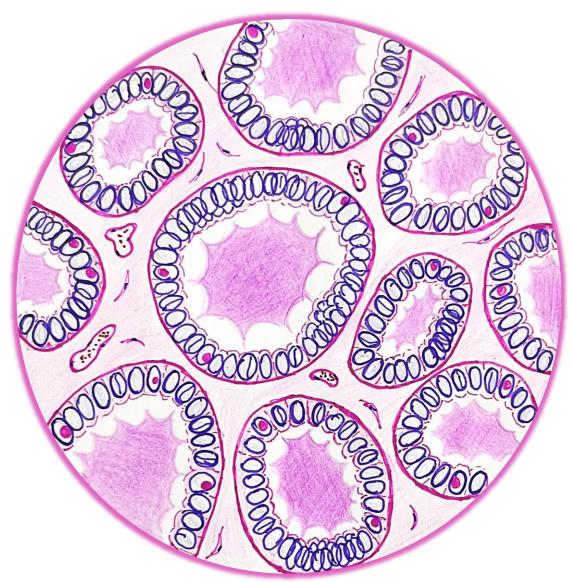
- Tumor has typical papillary architecture and is surrounded by a fibrous capsule
- Accounts for 10% of cases of PTC
- Prognosis excellent
- Regional lymph node metastasis may occur





Follicular variant

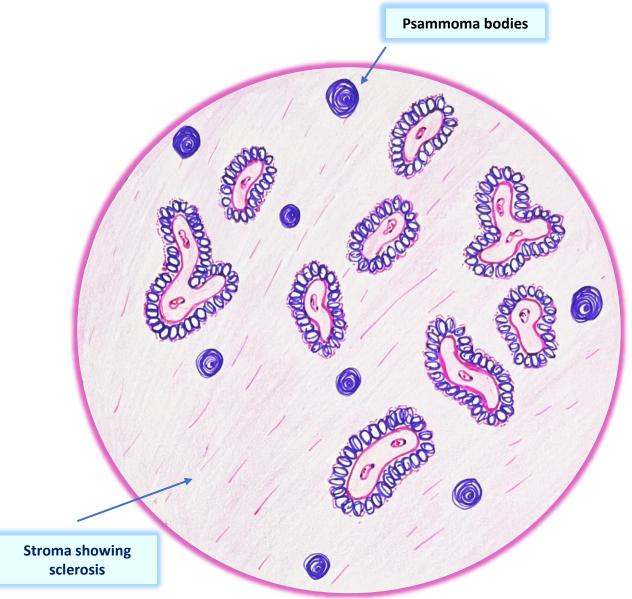
- Tumor has follicular growth pattern and can be infiltrative or encapsulated with invasion
- Nuclei of the cells lining the follicles have features of conventional PTC
- Colloid is thick appearing homogenous and strongly eosinophilic
- In some tumors, follicles are large cystically dilated – macrofollicular variant





Diffuse sclerosing variant

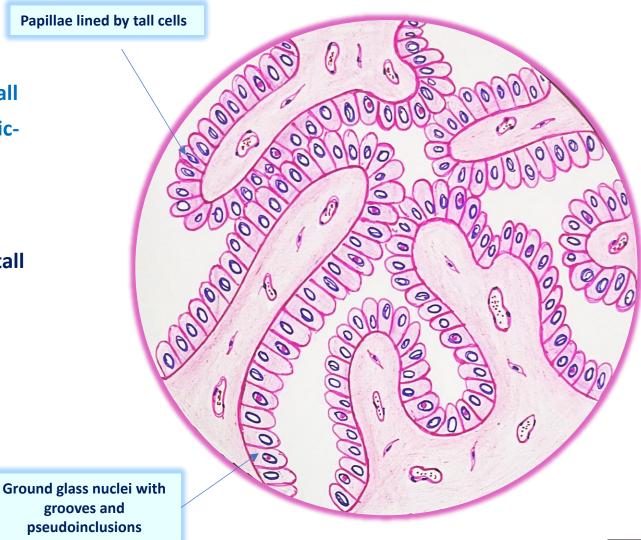
- Uncommon variant occurring in second and third decades and is common in women
- Clinical presentation diffuse enlargement of thyroid
- Microscopically entire thyroid or single lobe may be involved
- Characterized by dense sclerosis, numerous psammoma bodies, and chronic lymphocytic thyroiditis changes in background
- This variant is associated with extrathyroidal extension, cervical lymphnode metastasis and metastasis to distant organs mainly to lungs





Tall cell variant

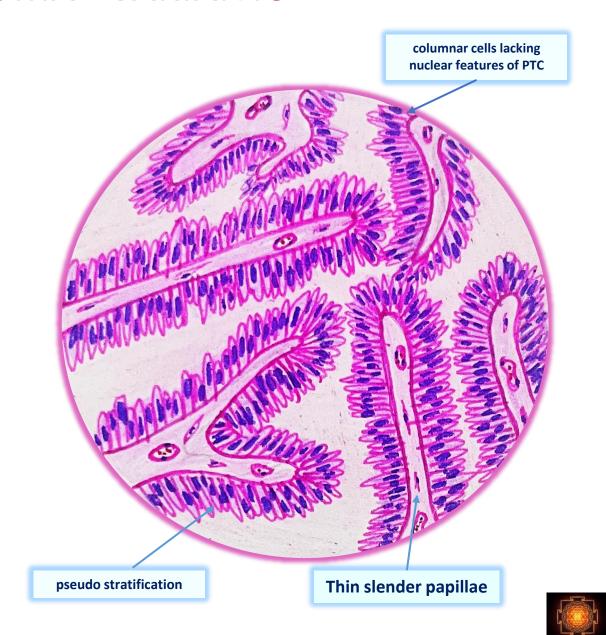
- Characterized by the cells which are two to three times as tall as they are wide and show abundant eosinophilic (oncocyticlike) cytoplasm
- Nuclear features of PTC are present
- Tall cells should account for > 30% of tumor to diagnose as tall cell variant
- Occurs in older individuals and has worse prognosis
- Extrathyroid extension and metastasis to other sites is common
- Prognosis- less favorable





Columnar cell variant

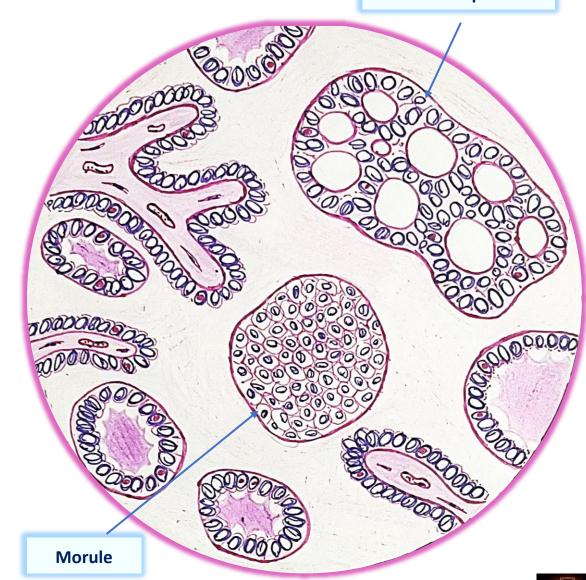
- Characterized by columnar cells with prominent pseudo stratification
- Cells lack nuclear features of conventional PTC
- These tumors are hypercellular neoplasms showing thin papillae or glandular like spaces lined by pseudostratified epithelium
- Tumor cells may show subnuclear vacuolization or clear cytoplasm similar to as seen in endometroid or intestinal adenocarcinoma



Cribriform pattern

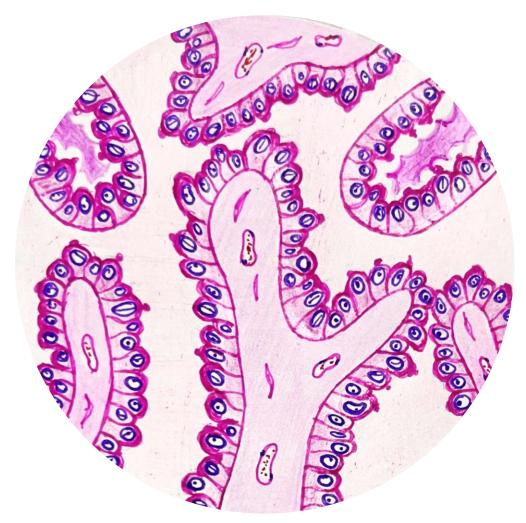
Cribriform or morular variant

- Occurs exclusively in females
- can occur sporadically or as a manifestation of familial adenomatous polyposis
- Morphology tumor is encapsulated and shows admixture of cribriform, trabecular, papillary, follicular and solid growth patterns with round squamoid structures called morules
- Papillae are lined by columnar shaped cells and intraluminal colloid is absent
- Nuclei may not be clear but have grooves and pseudoinclusions
- Morules contain optically clear nuclei



Hobnail variant

- Tumor should contain > 30% of cells with hobnail features
- Tumor shows complex papillary and micropapillary structures lined by follicular cells containing eosinophilic cytoplasm, apically located nuclei showing decreased N:C ratio and prominent nucleoli and has loss of cellular cohesion
- Extrathyroidal extension, angiolymphatic invasion, necrosis and atypical mitotic figures are common





Oncocytic variant

- Rare variant
- characterized by papillary tumors with oncocytic cell
- Cells have abundant eosinophilic granular cytoplasm
- Nuclear features are of papillary thyroid carcinoma





Solid or trabecular variant

- Characterized by solid or trabecular growth pattern
- Solid variant constitutes 1 3% of adult papillary carcinoma
- Tumor cells show features of conventional PTC
- Poor prognosis and is associated with lung metastasis



Other rare variants include

- Spindle cell variant
- Clear cell variant
- Warthin-like variant



Mode of spread

- Most common Lymphatic spread
- Minority of cases hematogenous spread to lungs

Diagnostic tests

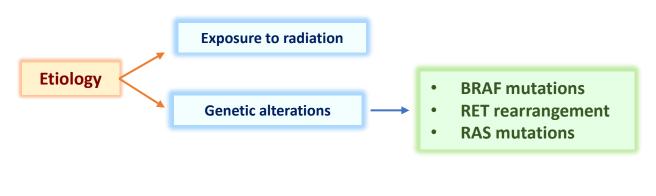
- Radionuclide scanning
- Fine needle aspiration cytology
- Scintigraphy presents as cold nodule
- Ultrasound hypoechoic or isoechoic nodule, microcalcifications and disorganized internal vascularity
- Biopsy



Prognosis

- Excellent with 10 year survival of about 95%
- Papillary microcarcinoma and Non-invasive follicular neoplasm with papillary like nuclear features excellent prognosis. Lobectomy is treatment (total thyroidectomy is not required)
- Local and regional recurrences 5% to 20% of patients
- Distant metastasis 10% to 15% of patients
- Less favourable prognostic factors are
 - Age more than 40 years
 - Presence of extrathyroidal extension
 - Presence of distant metastasis

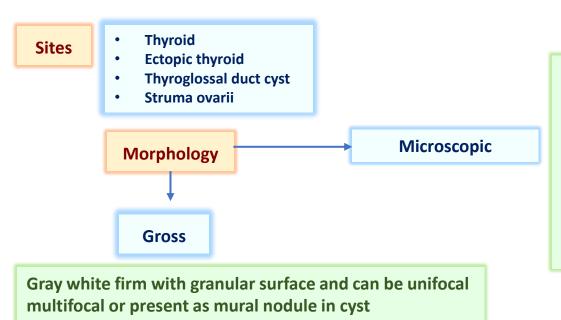




Precursor lesions

- Micropapillary for conventional PTC
- Non-invasive follicular thyroid neoplasm with papillary – like nuclear features- for follicular variant of PTC

Sex/age Female predominance in the age group of children to adults



- Papillae with fibrovascular core
- Characteristic nuclear features
 - Orphan annie eyed nuclei
 - Overlapping
 - Nuclear grooves
 - Nuclear pseudoincusion
- Psammoma bodies

Variants of papillary carcinoma

- Follicular variant
- Micropapillary
- **Encapsulated**
- Tall cell variant
- Columnar cell variant
- Hobnail variant
- Cribriform variant
- Oncocytic variant
- Solid/ trabecular
- Spindle cell variant
- Clear cell variant
- Warthin like



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